60060.....015563'' and in the table in paragraph (c)(2) in the entry for ''015563'' by removing the sponsor name ''Agribusiness Marketers, Inc.,'' and adding in its place ''Mallinckrodt Veterinary Operations, Inc., 421 East Hawley St., Mundelein, IL 60060.''

Dated: April 4, 1996.

Robert C. Livingston,

Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine.

[FR Doc. 96–9779 Filed 4–19–96; 8:45 am]

BILLING CODE 4160–01–F

21 CFR Part 558

New Animal Drugs; Change of Sponsor

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect the change of sponsor name for a new animal drug application (NADA) from MAC–PAGE, Inc., to ADM Animal Health & Nutrition Div.

EFFECTIVE DATE: April 22, 1996.

FOR FURTHER INFORMATION CONTACT: Thomas J. McKay, Center for Veterinary Medicine (HFV–102), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0213.

SUPPLEMENTARY INFORMATION: In the Federal Register of October 6, 1994 (59 FR 50828), FDA published a final rule amending the animal drug regulations to reflect the change of sponsors for all NADA's held by Central Soya, P. O. Box 1400, Fort Wayne, IN 46801–2508, including MAC–PAGE, Inc., 1600 South Wilson Ave., Dunn, NC 28334, a wholly-owned subsidiary of Central Soya, transferred to Premiere Agri Technologies, Inc. The subsidiaries retained their names and drug labeler codes. In the Federal Register of September 11, 1995 (60 FR 40752), FDA published a final rule amending the animal drug regulations to reflect the change of sponsor name from Premiere Agri Technologies, Inc., and the names of all wholly-owned subsidiaries, to ADM Animal Health & Nutrition Div., P.O. Box 2508, Fort Wayne, IN 46801–2508. Both final rules, which reflected these changes, inadvertently did not include NADA 131–957 (Tylosin). This document corrects that error. Accordingly, FDA is amending the regulations in 21 CFR 558.625 to reflect the change of sponsor.

List of Subject in 21 CFR Part 558

Animal drugs.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Center for Veterinary Medicine, 21 CFR part 558 is amended as follows:

PART 558—NEW ANIMAL DRUGS FOR USE IN ANIMAL FEEDS

1. The authority citation for 21 CFR part 558 continues to read as follows:

Authority: Secs. 512, 701 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b, 371).

§ 558.625 [Amended]

2. Section 558.625 *Tylosin* is amended in paragraph (b)(79) by removing ''047427'' and adding in its place ''012286''.

Dated: April 4, 1996.

Robert C. Livingston,

Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine.

[FR Doc. 96–9784 Filed 4–19–96; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

21 CFR Parts 1313 and 1316

[DEA No. 112C]

Implementation of the Domestic Chemical Diversion Control Act of 1993 (Pub. L. 103–200); Correction

AGENCY: Drug Enforcement Administration (DEA), Justice.

ACTION: Correction to final regulations.

SUMMARY: This document contains corrections to the final regulations which were published on Thursday, June 22, 1995 (60 FR 32447). The regulations related to the registration, recordkeeping and reporting requirements for manufacturers, distributors, importers and exporters of listed chemicals.

EFFECTIVE DATE: April 22, 1996.

FOR FURTHER INFORMATION CONTACT: G. Thomas Gitchel, Chief, Liaison and Policy Section, Office of Diversion Control, Drug Enforcement Administration, Washington, D.C. 20537, Telephone (202) 307–7297.

SUPPLEMENTARY INFORMATION: The final regulations that are the subject of these corrections implement the Domestic Chemical Diversion Control Act of 1993 (Pub. L. 103–200) (DCDCA). The regulations amend Title 21, Code of Federal Regulations, to add a new Part 1309 and revise certain sections in Parts 1310, 1313 and 1316. As published, the final regulations contain errors that could cause confusion in the regulated industry.

Accordingly, the publication June 22, 1995 of the final regulations to implement the DCDCA, which were the subject of Federal Register Document 95–14978, is corrected as follows:

PART 1313—[CORRECTED]

1. On page 32465, in the first column, the section heading which reads ''§ 1312.32 Requirement of authorization for international transactions.'' is corrected to read ''§ 1313.32 Requirement of authorization for international transactions.''

PART 1316—[CORRECTED]

2. On page 32465, in the third column, amendment Number 1 immediately following PART 1316—[AMENDED] is corrected to read as follows:

1. The authority citation for Part 1316, Subpart A is amended to read as follows:

Authority: 21 U.S.C. 822(f), 830(a), 871(b), 880, 958(f), 965.

Dated: April 16, 1996.

Stephen H. Greene,

Deputy Administrator, Drug Enforcement Administration.

[FR Doc. 96–9813 Filed 4–19–96; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF TRANSPORTATION

Federal Highway Administration

23 CFR Part 625

[FHWA Docket No. 95–12]

RIN 2125–AD38

Design Standards for Highways; Geometric Design of Highways and Streets

AGENCY: Federal Highway Administration (FHWA), DOT.

ACTION: Interim final rule; request for comments.

SUMMARY: The National Highway System (NHS) was established by the National Highway System Designation Act of 1995 (Pub. L. 104–59, 109 Stat. 568). To reflect the establishment of the NHS, the FHWA is revising several areas of the text in its regulation governing design standards for highways; updating the listing of standards; relocating the guides and references; and adopting as its interim